wherein R⁵ and R⁶ are each independently selected from the group consisting of -OC(O)OCH₃, -OH, -SH, -NH₂, -OSO₃H, -OPO₃H, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted monosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer, provided that at least one of R⁵ and R⁶ are -OC(O)OCH₃; wherein R⁷, R⁸, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶, R¹⁷, R¹⁸ and R¹⁹ are each

independently selected from the group consisting of -H, -OH, -SH, $-NH_2$, $-OSO_3H$, $-OPO_3H$, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and \mathbb{R}^7 and \mathbb{R}^8 together, \mathbb{R}^{12} and \mathbb{R}^{13} together, \mathbb{R}^{14} and \mathbb{R}^{15}

together, R^{16} and R^{17} together, and R^{18} and R^{19} together independently form a double bond to a moiety selected from the group consisting of =0, =S, =CH₂ and =NOH, provided that only one each of R^{12} and R^{13} or R^{18} and R^{19} can independently be -H; wherein R^{24} and R^{25} are either -H or -CH₃; wherein the dotted line is an optional double bond; wherein the -OC(0)OCH₃ at the 3 position is in either the α or β configuration; and a pharmaceutically acceptable excipient.

34. (New) The pharmaceutical composition of claim 33, wherein said at least one compound has the following structure

wherein R⁷, R⁸, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶, R¹⁷, R¹⁸ and R¹⁹ are each independently selected from the group consisting of -H, -OH, -SH, -NH₂, -OSO₃H, -OPO₃H, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted oligosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and R⁷ and R⁸ together, R¹² and R¹³ together, R¹⁴ and R¹⁵

together, R^{16} and R^{17} together, and R^{18} and R^{19} together independently form a double bond to a moiety selected from the group consisting of =0, =S, =CH₂ and =NOH, provided that only one each of R^{12} and R^{13} or R^{18} and R^{19} can independently be -H; wherein R^{24} and R^{25} are either -H or -CH₃; wherein the dotted line is an optional double bond; wherein the -OC(O)OCH₃ at the 3 position is in either the α or β configuration; and a pharmaceutically acceptable excipient.

35. (New) The pharmaceutical composition of claim 34, wherein said at least one compound has the following structure

wherein R⁷, R⁸, R¹², R¹³, R¹⁴, R¹⁵, R¹⁶ and R¹⁷ are each independently selected from the group consisting of -H, -OH, -SH, -NH₂, -OSO₃H, -OPO₃H, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heterocycle, an optionally substituted heterocycle, an optionally substituted monosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and R⁷ and R⁸ together, R¹² and R¹³ together, R¹⁴ and R¹⁵ together, and R¹⁶ and R¹⁷ together independently form a double bond to a moiety

selected from the group consisting of =0, =S, =CH2 and =NOH, provided that only one of each of R12 and R13 can independently be -H;

wherein R^{24} and R^{25} are either -H or -CH₃; wherein the dotted line is an optional double bond; wherein the $-OC(0)OCH_3$ at the 3 position is in either the α or β configuration;

and a pharmaceutically acceptable excipient.

36. (New) The pharmaceutical composition of claim 35, wherein said at least one compound has the following structure

wherein R¹² and R¹³ are each independently selected from the group consisting of -H, -OH, -SH, -NH₂, -OSO₃H, -OPO₃H, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted monosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and R^{12} and R^{13} together form a double bond to a moiety selected from the group consisting of =0, =S, = CH_2 and =NOH, provided that only one of R^{12} and R^{13} is -H;

wherein R^{24} and R^{25} are either -H or -CH₃: wherein the dotted line is an optional double bond; wherein the -OC(0)OCH $_3$ at the 3 position is in either the α or β configuration; and a pharmaceutically acceptable excipient.

37. (New) The pharmaceutical composition of claim 34, wherein said at least one compound has the following structure

$$R^{18}$$
 R^{24}
 R^{18}
 R^{19}
 R^{16}
 R^{17}
 R^{17}
 R^{17}
 R^{18}
 R^{18}
 R^{19}
 R^{16}
 R^{17}

wherein R^7 , R^8 , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} and R^{19} are each independently selected from the group consisting of -H, -OH, -SH, -NH2, -OSO3H, -OPO₃H, an ester, a phosphoester, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted monosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and R⁷ and R^8 together, R^{14} and R^{15} together, R^{16} and R^{17} together, and R^{18} and R¹⁹ together independently form a double bond to a moiety selected from the group consisting of =0, =S, =CH2 and =NOH, provided that only one of each of R18 and R19 can be -H; wherein R^{24} and R^{25} are either -H or -CH₃;

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wherein the dotted line is an optional double bond;

wherein the -OC(0)OCH $_3$ at the 3 position is in either the α or β configuration;

and a pharmaceutically acceptable excipient.

38. (New) The pharmaceutical composition of claim 37, wherein said at least one compound has the following structure

wherein R^{18} and R^{19} are each independently selected from the group consisting of -H, -OH, -SH, -NH₂,-OSO₃H, -OPO₃H, an ester, a phosphoseter, a phosphonoester, a sulfite ester, a sulfate ester, a thioester, an amide, a sulfonamide, an amino acid, an ether, a thioether, an acyl group, a carbonate, a carbamate, a sulfonamide, a halogen, an optionally substituted alkyl group, an optionally substituted alkenyl group, an optionally substituted alkynyl group, an optionally substituted aryl moiety, an optionally substituted heterocycle, an optionally substituted heteroaryl moiety, an optionally substituted monosaccharide, an optionally substituted oligosaccharide, a nucleoside, a nucleotide, an oligonucleotide, a polymer and R^{18} and R^{19} together form a double bond to a moiety selected from the group consisting of =O, =S, =CH₂ and =NOH, provided that only one of R^{18} and R^{19} is -H;

wherein R^{24} and R^{25} are either -H or -CH_3; wherein the dotted line is an optional double bond; wherein the -OC(O)OCH_3 at the 3 position is in either the α or β configuration;

and a pharmaceutically acceptable excipient.

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39. (New) The pharmaceutical composition of claim 34, wherein said at least one compound has the following structure

and a pharmaceutically acceptable excipient.

- 40. (New) A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof, comprising administering to the subject, or delivering to the subject's tissues, an effective amount of the pharmaceutical composition of claim 33.
- 41. (New) The method of claim 40, wherein the androgen responsive disease is selected from the group consisting of prostate cancer, benign prostatic hyperplasia, breast cancer, alopecia, acne, hypogonadism and hirsutism.
- 42. (New) A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof, comprising administering to the subject, or delivering to the subject's tissues, an effective amount of the pharmaceutical composition of claim 34.
- 43. (New) The method of claim 42, wherein the androgen responsive disease is selected from the group consisting of prostate cancer, benign prostatic hyperplasia, breast cancer, alopecia, acne, hypogonadism and hirsutism.

- 44. (New) A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof, comprising administering to the subject, or delivering to the subject's tissues, an effective amount of the pharmaceutical composition of claim 35.
- 45. (New) The method of claim 44, wherein the androgen responsive disease is selected from the group consisting of prostate cancer, benign prostatic hyperplasia, breast cancer, alopecia, acne, hypogonadism and hirsutism.
- 46. (New) A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof, comprising administering to the subject, or delivering to the subject's tissues, an effective amount of the pharmaceutical composition of claim 36.
- 47. (New) The method of claim 46, wherein the androgen responsive disease is selected from the group consisting of prostate cancer, benign prostatic hyperplasia, breast cancer, alopecia, acne, hypogonadism and hirsutism.
- 48. (New) A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof, comprising administering to the subject, or delivering to the subject's tissues, an effective amount of the pharmaceutical composition of claim 37.
- 49. (New) The method of claim 48, wherein the androgen responsive disease is selected from the group consisting of prostate cancer, benign prostatic hyperplasia, breast cancer, alopecia, acne, hypogonadism and hirsutism.
- 50. A method to treat or prevent an androgen responsive disease in a subject, or to ameliorate one or more symptoms thereof,

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